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OK protein - protein search, using sw model

Run on: June 21, 2002, 08:23:32 ; Search time 93.48 Seconds
(without alignments)
99.810 Million cell updates/sec

Title: US-09-351-778A-12

Sequence: 1 MTGSTIAPTIDYNTATATGL.....NEKIHRLDGKPCSLLLQYD 84

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 747574 seqs, 11073796 residues

Word size : 0

Total number of hits satisfying chosen parameters: 747574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

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21: /SIDSL/gcgdata/hold-geneseq/geneeqp-emb1/AA2000.DAT:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	84	100.0	84	22	AA61872
2	56	66.7	95	22	AA61868
3	56	66.7	101	19	AA59925
4	56	66.7	101	19	AAW78902
5	56	66.7	101	19	AAW5787
6	56	66.7	101	19	AA61187
7	56	66.7	101	19	AAW8003
8	56	66.7	101	21	AAW4407
9	56	66.7	101	22	AAW47591
10	56	66.7	101	22	AAW50206
11	56	66.7	101	22	AA61866

12	42	50.0	42	22	AA61876
13	33	39.3	78	22	AA61869
14	30	35.7	87	22	AA61870
15	28	33.3	40	22	AA61873
16	28	33.3	77	22	AA61871
17	19	22.6	93	22	AA61867
18	14	16.7	19	22	AA61874
19	13	15.5	94	22	AA61865
20	8	9.5	8	22	AA61875
21	7	8.3	157	21	AA61874
22	7	8.3	197	20	AA61853
23	7	8.3	242	21	AA604903
24	7	8.3	242	21	AA659416
25	7	8.3	273	22	AB605846
26	7	8.3	316	21	AA604902
27	7	8.3	316	21	AA659415
28	7	8.3	404	22	AB65340
29	7	8.3	473	22	AB626845
30	7	8.3	482	21	AA656786
31	7	8.3	604	22	AB67946
32	7	8.3	635	22	AB66264
33	6	7.1	15	22	AA678731
34	6	7.1	43	22	AB64015
35	6	7.1	43	22	AA655030
36	6	7.1	43	22	AAW7745
37	6	7.1	43	22	AAW21656
38	6	7.1	43	22	AAW37959
39	6	7.1	50	22	AA678805
40	6	7.1	50	22	AA678832
41	6	7.1	51	22	AA678806
42	6	7.1	51	22	AA678831
43	6	7.1	52	22	AAU8951
44	6	7.1	52	22	AA678807
45	6	7.1	52	22	AA678830

ALIGNMENTS

RESULT 1
ID AAB61872 standard; Protein: 84 AA.
XX
AC AAB61872;
XX
DT 08-MAY-2001 (first entry)
XX
DE Ad2 ADP mutant d1737.
XX
KW Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;
anti-cancer; gene therapy; cytosolic; Ad2; mutant.
XX
OS Mastadenovirus.
XX
PN W0200104282-A2.
XX
PD 18-JAN-2001.
XX
PF 12-JUL-2000; 2000MO-US18971.
XX
PR 12-JUL-1999; 9905-0351778.
XX
PA (USL-) UNIV SAINT LOUIS.
XX
PI W01d WSM, Toth K, Doronin K, Tollefson AE;
XX
DR WPI: 2001-103079/11.
XX
PT Recombinant vector which is replication-competent in a neoplastic cell
PT and overexpresses an adenovirus death protein, useful in cancer therapy
PT when used together with replication-defective adenovirus which
PT expresses an anti-cancer gene -
XX

PS Example 9; Fig 20; 196pp; English.

XX The invention relates to a recombinant vector (V1) which is replication-
 CC competent in a neoplastic cell and which overexpresses an adenovirus
 CC death protein (ADP). The vector can be used in a method for promoting
 CC death of a neoplastic cell that comprises contacting the neoplastic cell
 CC with at least one V1; and a composition comprising V1 and a second
 CC recombinant virus which is: (a) replication defective and which
 CC expresses an anti-cancer gene product, where V1 complements replication
 CC of the second recombinant virus; or (b) replication-competent in a
 CC neoplastic cell. V1, together with one or more replication-defective
 CC adenovirus which expresses an anti-cancer gene product, are useful in
 CC cancer therapy. Overexpression of ADP by V1 results in faster lysis of
 CC cells and spread of the virus throughout a cell monolayer than viruses
 CC expressing wild-type levels of ADP. The present sequence represents the
 CC amino acid sequence of an Ad2 ADP mutant.

XX Sequence 84 AA:

SO

Query Match 100.0%; Score 84; DB 22; Length 84;
 Best Local Similarity 100.0%; Pred. No. 5; 9e-79;
 Matches 84; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MTGCTIAPTDDYNTATGCTGATLALNPQIALMFVCLIMMLICLKRARRAPPIYRPIIV 60
 DB 1 mtgcttaptddyntatgctgatsalnlpqialmfvcclimmlcclkrarrappiyrrpiiv 60
 OY 61 LNPHEKIHRLDGLKPCSLLOYD 84
 DB 61 lnphekihrlldglkpcslloyd 84

DB 61 lnphekihrlldglkpcslloyd 84

RESULT 2
 AAB61868
 ID AAB61868 standard; Protein: 95 AA.

XX AAB61868:
 XX 08-MAY-2001 (first entry)
 XX
 XX Ad6 encoded adenovirus death protein (ADP).
 XX
 XX Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;
 XX anti-cancer; gene therapy; cytostatic; Ad6.
 XX Mastadenovirus.
 XX
 XX OS
 XX
 XX Key Location/Qualifiers
 XX Peptide 1..26
 XX /note- "Fragment specifically claimed for"
 XX Peptide 41..59
 XX /note- "Fragment specifically claimed for"
 XX Peptide 63..70
 XX /note- "Fragment specifically claimed for"

XX WO200104282-A2.
 XX
 XX 18-JAN-2001.
 XX
 XX 12-JUL-2000; 2000WO-US18971.
 XX
 XX 12-JUL-1999; 99US-0351778.
 XX
 XX (URSL-) UNIV SAINT LOUIS.
 XX
 XX Wold WSM, Toth K, Doronin K, Tollefson AE;
 XX
 XX WPI; 2001-103079/11.
 XX
 XX Recombinant vector which is replication-competent in a neoplastic cell
 XX PT and overexpresses an adenovirus death protein, useful in cancer therapy
 XX when used together with replication-defective adenovirus which

PT expresses an anti-cancer gene -

PS Claim 5; Page 157; 196pp; English.

XX The invention relates to a recombinant vector (V1) which is replication-
 CC competent in a neoplastic cell and which overexpresses an adenovirus
 CC death protein (ADP). The vector can be used in a method for promoting
 CC death of a neoplastic cell that comprises contacting the neoplastic cell
 CC with at least one V1; and a composition comprising V1 and a second
 CC recombinant virus which is: (a) replication defective and which
 CC expresses an anti-cancer gene product, where V1 complements replication
 CC of the second recombinant virus; or (b) replication-competent in a
 CC neoplastic cell. V1, together with one or more replication-defective
 CC adenovirus which expresses an anti-cancer gene product, are useful in
 CC cancer therapy. Overexpression of ADP by V1 results in faster lysis of
 CC cells and spread of the virus throughout a cell monolayer than viruses
 CC expressing wild-type levels of ADP. The present sequence represents the
 CC amino acid sequence of an ADP encoded by Ad6.

XX Sequence 95 AA:

SO

Query Match 66.7%; Score 56; DB 22; Length 95;
 Best Local Similarity 100.0%; Pred. No. 4; 9e-50;
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMFVCLIMMLICLKRARRAPPIYRPIIVLNPHEKIHRLDGLKPCSLLOYD 84
 DB 40 ialmfvcclimmlcclkrarrappiyrrpiivlnphekihrlldglkpcslloyd 95

RESULT 3
 AAM59925
 ID AAM59925 standard; Protein: 101 AA.

XX AAM59925:
 XX 11-JAN-1999 (first entry)
 XX
 XX Adenovirus death protein.
 XX
 XX Adenovirus death protein; ADP; transcription regulatory element;
 XX vector; breast cancer; prostate cancer; liver cancer; colon cancer;
 XX gene therapy.
 XX Mastadenovirus.
 XX
 XX OS
 XX
 XX WO9839464-A2.
 XX
 XX 11-SEP-1998.
 XX
 XX 03-MAR-1998; 98WO-US04080.
 XX
 XX 02-MAR-1998; 98US-0054523.
 XX 03-MAR-1997; 97US-0039762.
 XX 03-MAR-1997; 97US-0039763.
 XX 04-AUG-1997; 97US-0054523.
 XX
 XX (CALY-) CALYDON INC.
 XX
 XX Henderson DR, Lamparski HG, Yu D;
 XX
 XX WPI; 1998-495860/42.
 XX N-PSDB; AAV53632.
 XX
 XX New adenovirus vectors, used for treating tumours - comprising first
 XX PT and second adenovirus genes under control of different heterologous
 XX PT transcriptional regulatory elements
 XX
 XX Disclosure; Page 94; 130pp; English.
 XX
 XX This is the amino acid sequence of adenovirus death protein (ADP).
 XX CC The invention provides replication-competent adenovirus vectors

CC specific for target cells and methods of using such vectors. The
CC vectors contain heterologous transcription regulatory elements
CC (TREs) and may incorporate a gene, such as the ADP gene (see
CC AAV53632), which can contribute to cytotoxicity in the target cell.
CC Adenoviral replication can be restricted to target cells in which
CC the heterologous TREs are functional and thus the vectors can
CC provide selective cytotoxicity to the target cells (e.g. prostate,
CC liver, breast or colon), particularly neoplastic cells.

XX Sequence 101 AA:

Query Match 66.7%; Score 56; DB 19; Length 101;
Best Local Similarity 100.0%; Pred. No. 5.2e-50;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMPVCLIIIMWLICCLRRARPPIYRIIYLNPHNKHIRLDGLKPCSLLOYD 84
DB 46 IALMFVCLIIIMWLICCLKRRARPPIYRIIYLNPHNKHIRLDGLKPCSLIYQD 101

RESULT 4

AAW78902 standard; Protein; 101 AA.

AAW78902:

21-DEC-1998 (first entry)

Adenovirus death protein.

Carcinoma embryonic antigen; transcriptional regulatory element;

CEA-TRE; human; promoter; enhancer; vector; cancer; gene therapy;

PCR; primer; adenovirus death protein; ADP.

Mastadenovirus.

MO9839467-A2.

11-SEP-1998.

03-MAR-1998; 98MO-US04133.

02-MAR-1998; 98US-0039763.

03-MAR-1997; 97US-0039763.

(CALY-) CALYDON INC.

Henderson DR, Lamparski HG, Schuur ER;

WPI: 1998-495862/42.

N-PSDB; AAV52966.

New adenovirus vectors, particularly for cancer therapy - comprising

adenovirus gene under transcriptional control of carcinoembryonic

antigen transcriptional regulatory element

Disclosure; Page 68; 95pp; English.

This is the amino acid sequence of adenovirus death protein (ADP).

Ad gene under transcriptional control of a CEA-TRE. The vectors can

be used to detect and monitor samples for the presence of cells that

allow a CEA-TRE to function, and to selectively kill such cells,

especially malignant cells. Vectors containing an ADP gene (see

AAV52966) may be more potent than vectors lacking the gene, making

possible more effective treatment and/or lower dosage requirement.

Sequence 101 AA:

Query Match 66.7%; Score 56; DB 19; Length 101;
Best Local Similarity 100.0%; Pred. No. 5.2e-50;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMPVCLIIIMWLICCLRRARPPIYRIIYLNPHNKHIRLDGLKPCSLLOYD 84
DB 46 IALMFVCLIIIMWLICCLKRRARPPIYRIIYLNPHNKHIRLDGLKPCSLIYQD 101

RESULT 5

AAW75787 standard; Protein; 101 AA.

AAW75787:

21-DEC-1998 (first entry)

Adenovirus death protein.

Probasin transcriptional response element; PB-TRE; rat;

androgen receptor; adenovirus; vector; prostate cancer;

gene therapy; adenovirus death protein; ADP.

Mastadenovirus.

MO9839466-A2.

11-SEP-1998.

03-MAR-1998; 98MO-US04132.

02-MAR-1998; 98US-0033333.

03-MAR-1997; 97US-0039762.

(CALY-) CALYDON INC.

Henderson DR, Lamparski HG, Schuur ER, Yu D;

WPI: 1998-506369/43.

N-PSDB; AAV57354.

New adenovirus vectors, particularly for cancer therapy - comprising

adenovirus gene under transcriptional control of a probasin

transcriptional regulatory element

Disclosure; Page 96; 117pp; English.

This is the amino acid sequence of adenovirus death protein (ADP).

Ad gene under transcriptional control of a probasin transcriptional

response element (PB-TRE, see AAV57354). The vector can be used for

detecting cells that allow a PB-TRE to function, especially cells

expressing an androgen receptor, such as prostate cells. They can

be used to confer selective toxicity to such cells. In particular,

the vectors can be used for treating cancers such as prostate cancer.

Ad vectors containing the ADP gene (see AAV57354) may render the

vector more potent, making possible more effective treatment and/or

a lower dosage requirement. An Ad vector has been constructed that

contains the ADP gene under control of PB-TRE. Cytotoxicity was

demonstrated toward LNCaP (prostate carcinoma) cells.

Sequence 101 AA:

Query Match 66.7%; Score 56; DB 19; Length 101;
Best Local Similarity 100.0%; Pred. No. 5.2e-50;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMPVCLIIIMWLICCLRRARPPIYRIIYLNPHNKHIRLDGLKPCSLLOYD 84
DB 46 IALMFVCLIIIMWLICCLKRRARPPIYRIIYLNPHNKHIRLDGLKPCSLIYQD 101

RESULT 6

AAW61197

ID AAM61197 standard; Protein: 101 AA.
 XX
 AC AAM61197;
 XX
 DT 07-DEC-1998 (first entry)
 XX
 DE Adenovirus death protein.
 XX
 KW Adenovirus death protein: ADP; vector: hepatoma; cancer;
 KM alpha-fetoprotein transcription regulatory element; AFP-TRE;
 KM hepatocellular carcinoma; hepatoma; gene therapy; human.
 XX
 OS Mastadenovirus type 2.
 XX
 PN MO9839465-A2.
 XX
 PD 11-SEP-1998.
 XX
 PF 03-MAR-1998; 98MO-US04084.
 XX
 PR 02-MAR-1998; 98US-0039597.
 PR 03-MAR-1997; 97US-0039597.
 XX
 PA (CALY-) CALYDON INC.
 XX
 PI Henderson DR, Lamparski HG, Little AS, Schuur ER;
 XX
 DR WPI: 1998-495861/42.
 DR N-PSDB: AAV47675.
 XX
 PT New adenovirus vector, for treating cancers - comprising an
 PT adenovirus gene under the transcriptional control of an alpha
 PT fetoeprotein transcription regulatory element
 XX
 PS Claim 29; Page 74; 102pp; English.
 XX
 CC This is the amino acid of the adenovirus death protein (ADP) of
 CC of adenovirus type 2. The ADP coding sequence (see AAV47675), with
 CC or without the 5' leader, can be introduced into an adenoviral
 CC genome, e.g. in the E3 or E4 region. Inclusion of such a coding
 CC sequence in an adenoviral vector significantly enhances the extent
 CC of cytotoxicity, cell killing and virus production. The invention
 CC provides replication-competent adenovirus vectors which
 CC preferentially replicate in cells that express alpha-fetoprotein
 CC (AFP), particularly hepatoma cells. The vectors comprise at
 CC least one adenovirus gene, preferably a gene that contributes to
 CC cytotoxicity, under the transcriptional control of an AFP
 CC transcription regulatory element (see AAV47654-55). The vectors
 CC are useful for conferring selective cytotoxicity to AFP-expressing
 CC cells, especially cancer cells.
 CC
 XX
 SQ Sequence 101 AA;

Query Match 66.7%; Score 56; DB 19; Length 101;
 Best Local Similarity 100.0%; Pred. No. 5.2e-50;
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMFVCLIIIMWLICCKRRRAPPIYPIVLPNHNKIRHLDGLKPCSLLOYD 84
 XX
 DB 46 IALMFVCLIIIMWLICCKRRRAPPIYPIVLPNHNKIRHLDGLKPCSLLOYD 101

RESULT 7
 ID AAM98003 standard; Protein: 101 AA.
 XX
 AC AAM98003;
 XX
 DT 21-JUN-1999 (first entry)
 XX
 DE Adenovirus death protein.
 XX

KW Enhancer; glandular kallikrein-1; hK2; human;
 KM prostate cancer; therapy; adenovirus death protein.
 XX
 OS Mastadenovirus 2.
 XX
 PN MO9906576-A1.
 XX
 PD 11-FEB-1999.
 XX
 PF 04-AUG-1998; 98MO-US16312.
 XX
 PR 03-AUG-1998; 98US-0127834.
 PR 04-AUG-1997; 97US-0054523.
 PR 02-MAR-1998; 98US-0076545.
 XX
 PA (CALY-) CALYDON INC.
 XX
 PI Henderson DR, Schuur ER, Yu D;
 XX
 DR WPI: 1999-153804/13.
 DR N-PSDB: AAX24756.
 XX
 PT New nucleic acid containing the human glandular kallikrein enhancer
 PT - providing increased expression of heterologous sequences in
 PT prostatic cells, and related adenoviral vectors for treating
 PT prostatic cancer
 XX
 PS Disclosure; Page 165-166; 179pp; English.
 XX
 CC This protein comprises the adenovirus death protein (ADP) of
 CC adenovirus serotype 2. The invention provides novel adenovirus
 CC vectors in which at least one adenovirus gene, preferably one that
 CC contributes to cytotoxicity, is placed under transcriptional
 CC control of a human glandular kallikrein hK2 enhancer
 CC CC transcriptional regulatory element (hK2-TRE; see AAX24755). Such
 CC vectors are useful for treatment of cancers such as prostate
 CC cancer. The ADP gene may render the adenoviral vector more potent,
 CC making possible more effective treatment and/or lower dosage
 CC requirement.
 CC
 XX
 SQ Sequence 101 AA;

Query Match 66.7%; Score 56; DB 20; Length 101;
 Best Local Similarity 100.0%; Pred. No. 5.2e-50;
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMFVCLIIIMWLICCKRRRAPPIYPIVLPNHNKIRHLDGLKPCSLLOYD 84
 XX
 DB 46 IALMFVCLIIIMWLICCKRRRAPPIYPIVLPNHNKIRHLDGLKPCSLLOYD 101

RESULT 8
 ID AAY84407 standard; Protein: 101 AA.
 XX
 AC AAY84407;
 XX
 DT 25-JUL-2000 (first entry)
 XX
 DE Amino acid sequence of an adenoviral death protein.
 XX
 KW Adenoviral vector; adenovirus gene; transcriptional control;
 KM transcriptional regulatory element; TRE; adenoviral propagation;
 KM death protein; tumour.
 XX
 OS Mastadenovirus.
 XX
 PN MO200015820-A1.
 XX
 PD 23-MAR-2000.
 XX
 PF 10-SEP-1999; 99MO-US20718.

XX 10-SEP-1998; 98US-0099791.
PR 09-SEP-1999; 99US-0099791.
XX (CALY-) CALYDON INC.
PA Yu DC, Henderson DR:
XX WPI: 2000-271456/23.
DR N-PSDB: AA299937.
PT Adenovirus vectors comprising cell-status specific response elements
XX useful in gene therapy protocols for the treatment of cancers -
PS Disclosure; Fig 9; 79pp; English.
XX The present sequence represents an adenoviral death protein, which is
CC used to construct the vectors of the invention. The specification
CC describes adenoviral vectors which comprise an adenovirus gene
CC under transcriptional control of a cell status specific transcriptional
CC regulatory element (TRE). The TRE is preferably one that is
CC essential for adenoviral propagation. The adenovirus vectors
CC may be used for the treatment of a range of tumours such as lung,
CC stomach, breast, colon and rectum, and uterine and cervix cancers.
XX Sequence 101 AA:

Query Match 66.7%; Score 56; DB 21; Length 101;
Best Local Similarity 100.0%; Pred. No. 5.2e-50;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALFVCLIIIMWLIICLKRBRAPPIYRPIIVLNPHEKIHRLDGAKPSILLOYD 84
DB 46 IALFVCLIIIMWLIICLKRBRAPPIYRPIIVLNPHEKIHRLDGAKPSILLOYD 101

RESULT 9
AAB47591
ID AAB47591 standard; Protein: 101 AA.
XX
AC AAB47591;
XX
DT 07-JAN-2002 (first entry)
XX
DE ADP amino acid sequence.
XX
KW Adenovirus; ADP: replication-competent; adenoviral vector; TRE;
KW transcriptional regulatory element; mutation; deletion; IRES;
KW promoter; internal ribosome entry site; cytotoxic; cancer; bladder.
XX
OS Adenovirus.
XX
PN WO200173093-A2.
XX
PD 04-OCT-2001.
XX
PF 21-MAR-2001; 2001WO-US09036.
XX
PR 24-MAR-2000; 2000US-192156P.
XX
PA (CALY-) CALYDON INC.
XX
PI Yu D, Li Y, Henderson DR:
XX
DR WPI: 2001-639234/73.
DR N-PSDB: AA43535.
XX
PT Replication-competent adenoviral vector, useful e.g. for killing cancer
PT cells, contains two genes linked by internal ribosome entry site and
PT controlled by target-specific regulator -
XX
PS Disclosure; Fig 9; 148pp; English.

XX This sequence represents adenoviral ADP. The ADP coding sequence may
CC be used in the replication-competent adenoviral vector (A) of the
CC invention which contains two genes (G1, G2) that are co-transcribed
CC as a single mRNA and under control of a heterologous, target cell-
CC specific transcriptional regulatory element (TRE). G2 has a mutation
CC in, or deletion of, its endogenous promoter and is controlled from
CC an internal ribosome entry site (IRES). The ADP coding sequence may
CC be used as G1 or G2. (A) has greater specificity for a target cell
CC than a similar vector in which TRE is operably linked to a gene and
CC which lacks an IRES. (A) are used to modify the genotype of target
CC cells, optionally in vitro with subsequent return of altered cells to
CC the host and where G2 is a cytotoxic gene, to confer selective
CC cytotoxicity to target cells, especially for killing cancer cells.
CC ADP displays a cytotoxic, particularly cell lysis, function. Also (A)
CC are used for diagnosis and monitoring, e.g. detection of bladder cancer
CC cells. The target cell-specific TRE ensures that (A) has better
CC targeting specificity, with minimal replication in non-target cells, so
CC a runaway infection is prevented but production of adenoviral proteins
CC in target cells activates and/or stimulates the immune response against
CC target cells producing such proteins. The use of an IRES (rather than
CC two identical control elements) eliminates the risk of homologous
CC recombination and may provide enough extra space for an additional
CC (therapeutic) gene.
XX Sequence 101 AA:

Query Match 66.7%; Score 56; DB 21; Length 101;
Best Local Similarity 100.0%; Pred. No. 5.2e-50;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALFVCLIIIMWLIICLKRBRAPPIYRPIIVLNPHEKIHRLDGAKPSILLOYD 84
DB 46 IALFVCLIIIMWLIICLKRBRAPPIYRPIIVLNPHEKIHRLDGAKPSILLOYD 101

RESULT 10
AAM50206
ID AAM50206 standard; Protein: 101 AA.
XX
AC AAM50206;
XX
DT 07-JAN-2002 (first entry)
XX
DE Adenovirus death protein.
XX
DE Adenovirus death protein.
XX
KW Adenovirus death protein; uroplakin II; vector;
KW transcriptional regulatory element; TRE; urothelial cell;
KW bladder cancer; human; gene therapy.
XX
OS Mastadenovirus 2.
XX
PN WO200172994-A2.
XX
PD 04-OCT-2001.
XX
PF 21-MAR-2001; 2001WO-US09224.
XX
PR 24-MAR-2000; 2000US-191861P.
XX
PA (CALY-) CALYDON INC.
XX
PI Yu D, Zhang H, Henderson DR:
XX
DR WPI: 2001-639229/73.
DR N-PSDB: AA170186.
XX
PT Human urothelial cell specific uroplakin transcriptional regulatory
PT sequences, useful for producing adenoviral vectors which can be used to
PT confer selective cytotoxicity to target cells, especially bladder
PT cancer cells -
XX

PS Example 6; Fig 12; 147pp; English.

XX The present sequence is that of the adenovirus death protein (ADP).

CC The ADP gene coding region (see A170186) was obtained by PCR

CC amplification and used in the construction of adenoviral vectors in

CC which ADP expression was under the control of a urothelial

CC cell-specific transcriptional regulatory element (TRE) derived from

CC the human uroplakin II gene 5' flanking region (see A170144). This

CC is an example of adenoviral vectors of the invention. Such vectors

CC comprise a gene, preferably an adenovirus gene, under transcriptional

CC control of a urothelial cell-specific TRE. They display urothelial

CC cell-specific cytotoxicity and are used for the specific, targeted

CC gene therapy of bladder cancer.

XX Sequence 101 AA:

SO

Query Match 66.7%; Score 56; DB 22; Length 101;

Best Local Similarity 100.0%; Pred. No. 5, 2e-50;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMFVCLIMWLICCLKRRRAPPYRPIIVLNPHNEKIHRLDGLKPSLLQYD 84

DB 46 IALMFVCLIMWLICCLKRRRAPPYRPIIVLNPHNEKIHRLDGLKPSLLQYD 101

RESULT 11

AAB61866

ID AAB61866 standard; Protein: 101 AA.

XX AAB61866;

AC

DT 08-MAY-2001 (first entry)

XX

DE Ad2 encoded adenovirus death protein (ADP).

XX

KW Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;

KM anti-cancer; gene therapy; cytostatic; Ad2.

XX

OS Mastadenovirus.

XX

OS

FH Key Location/Qualifiers

FT Peptide 1..26

FT /note= "fragment specifically claimed for"

FT Domain 1..40

FT /note= "putative luminal domain (AAB61873)"

FT Domain 41..59

FT /note= "transmembrane domain (AAB61874);"

FT Domain 63..70

FT /note= "fragment specifically claimed for"

FT Domain 71..70

FT /note= "cytosolic basic proline domain (AAB61875)"

FT Domain 60..101

FT /note= "fragment specifically claimed for"

FT Domain 60..101

FT /note= "cytoplasmic-nucleoplasmic domain"

XX WO200104282-A2.

XX

XX 18-JAN-2001.

XX

XX 12-JUL-2000; 2000MO-US18971.

XX

XX 12-JUL-1999; 99US-0351778.

XX

XX (UWSL-) UNIV SAINT LOUIS.

XX

XX Wold WSM, Toch K, Doronin K, Tollefson AE;

XX

XX WPI: 2001-103079/11.

XX

PT Recombinant vector which is replication-competent in a neoplastic cell

PT and overexpresses an adenovirus death protein, useful in cancer therapy

PT when used together with replication-defective adenovirus which

PT expresses an anti-cancer gene.

XX Claim 5; Page 156; 196pp; English.

XX

XX The invention relates to a recombinant vector (V1) which is replication-

CC competent in a neoplastic cell and which overexpresses an adenovirus

CC death protein (ADP). The vector can be used in a method for promoting

CC death of a neoplastic cell that comprises contacting the neoplastic cell

CC with at least one V1; and a composition comprising V1 and a second

CC recombinant virus which is: (a) replication defective and which

CC expresses an anti-cancer gene product, where V1 complements replication

CC of the second recombinant virus; or (b) replication-competent in a

CC neoplastic cell. V1, together with one or more replication-defective

CC adenovirus which expresses an anti-cancer gene product, are useful in

CC cancer therapy. Overexpression of ADP by V1 results in faster lysis of

CC cells and spread of the virus throughout a cell monolayer than viruses

CC expressing wild-type levels of ADP. The present sequence represents the

CC amino acid sequence of an ADP encoded by Ad2.

XX

SO Sequence 101 AA:

Query Match 66.7%; Score 56; DB 22; Length 101;

Best Local Similarity 100.0%; Pred. No. 5, 2e-50;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 29 IALMFVCLIMWLICCLKRRRAPPYRPIIVLNPHNEKIHRLDGLKPSLLQYD 84

DB 46 IALMFVCLIMWLICCLKRRRAPPYRPIIVLNPHNEKIHRLDGLKPSLLQYD 101

RESULT 12

AAB61876

ID AAB61876 standard; Peptide: 42 AA.

XX AAB61876;

AC

DT 08-MAY-2001 (first entry)

XX

DE Ad2 ADP cytosolic domain fragment.

XX

KW Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;

KM anti-cancer; gene therapy; cytostatic; Ad2.

XX

OS Mastadenovirus.

XX

OS

FH Key Location/Qualifiers

FT Peptide 1..26

FT /note= "fragment specifically claimed for"

FT Domain 1..40

FT /note= "putative luminal domain (AAB61873)"

FT Domain 41..59

FT /note= "transmembrane domain (AAB61874);"

FT Domain 63..70

FT /note= "fragment specifically claimed for"

FT Domain 71..70

FT /note= "cytosolic basic proline domain (AAB61875)"

FT Domain 60..101

FT /note= "fragment specifically claimed for"

FT Domain 60..101

FT /note= "cytoplasmic-nucleoplasmic domain"

XX WO200104282-A2.

XX

XX 18-JAN-2001.

XX

XX 12-JUL-2000; 2000MO-US18971.

XX

XX 12-JUL-1999; 99US-0351778.

XX

XX (UWSL-) UNIV SAINT LOUIS.

XX

XX Wold WSM, Toch K, Doronin K, Tollefson AE;

XX

XX WPI: 2001-103079/11.

XX

PT Recombinant vector which is replication-competent in a neoplastic cell

PT and overexpresses an adenovirus death protein, useful in cancer therapy

PT when used together with replication-defective adenovirus which

PT expresses an anti-cancer gene.

XX

XX Example 9; Fig 20; 196pp; English.

XX

XX The invention relates to a recombinant vector (V1) which is replication-

CC competent in a neoplastic cell and which overexpresses an adenovirus

CC death protein (ADP). The vector can be used in a method for promoting

CC death of a neoplastic cell that comprises contacting the neoplastic cell

CC with at least one V1; and a composition comprising V1 and a second

CC recombinant virus which is: (a) replication defective and which

CC expresses an anti-cancer gene product, where V1 complements replication

CC of the second recombinant virus; or (b) replication-competent in a
CC neoplastic cell. VI, together with one or more replication-defective
CC adenovirus which expresses an anti-cancer gene product, are useful in
CC cancer therapy. Overexpression of ADP by VI results in faster lysis of
CC cells and spread of the virus throughout a cell monolayer than viruses
CC expressing wild-type levels of ADP. The present sequence represents the
CC amino acid sequence of an Ad2 ADP cytosolic domain fragment.

XX Sequence 42 AA:

Query Match 50.0%; Score 42; DB 22; Length 42;

Best Local Similarity 100.0%; Pred. No. 6,5e-36; Mismatches 0; Indels 0; Gaps 0;

YY 43 CCLKRRRAPPYRPIVLPNPHNEKHRLDGLKPCSLLDQYD 84
DB 1 cclkrtrrrpplyrplvlvlnphnekhlrdglkpcslldqyd 42

RESULT 13

AA61869 standard; Protein: 78 AA.

XX AAB61869;

XX 08-MAY-2001 (first entry)

XX Ad2 ADP mutant dl716.

XX Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;

XX anti-cancer; gene therapy; cytostatic; Ad2; mutant.

XX Mastadenovirus.

XX MO200104282-A2.

XX 18-JAN-2001.

XX 12-JUL-2000: 2000MO-US18971.

XX 12-JUL-1999: 99US-0351778.

XX (UYSL-) UNIV SAINT LOUIS.

XX MOLD MSM, Toch K, Doronin K, Tollefson AE;

XX WPI: 2001-103079/11.

XX Recombinant vector which is replication-competent in a neoplastic cell

XX and overexpresses an adenovirus death protein, useful in cancer therapy

XX when used together with replication-defective adenovirus which

XX expresses an anti-cancer gene -

XX Example 9; Fig 20: 196pp; English.

XX The invention relates to a recombinant vector (VI) which is replication-

XX competent in a neoplastic cell and which overexpresses an adenovirus

XX death protein (ADP). The vector can be used in a method for promoting

XX death of a neoplastic cell that comprises contacting the neoplastic cell

XX with at least one VI; and a composition comprising VI and a second

XX recombinant virus which is: (a) replication defective and which

XX expresses an anti-cancer gene product, where VI complements replication

XX of the second recombinant virus; or (b) replication-competent in a

XX neoplastic cell. VI, together with one or more replication-defective

XX adenovirus which expresses an anti-cancer gene product, are useful in

XX cancer therapy. Overexpression of ADP by VI results in faster lysis of

XX cells and spread of the virus throughout a cell monolayer than viruses

XX expressing wild-type levels of ADP. The present sequence represents the

Query Match 39.3%; Score 33; DB 22; Length 78;
Best Local Similarity 100.0%; Pred. No. 2.1e-26;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

YY 29 IALMFVCLIMMVICLKRRRAPPYRPIVL 61
DB 46 ialmfvcilimviclckrrrrapplyrplvl 78

RESULT 14

AA61870 standard; Protein: 87 AA.

XX AAB61870;

XX 08-MAY-2001 (first entry)

XX Ad2 ADP mutant dl715.

XX Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;

XX anti-cancer; gene therapy; cytostatic; Ad2; mutant.

XX Mastadenovirus.

XX MO200104282-A2.

XX 18-JAN-2001.

XX 12-JUL-2000: 2000MO-US18971.

XX 12-JUL-1999: 99US-0351778.

XX (UYSL-) UNIV SAINT LOUIS.

XX MOLD MSM, Toch K, Doronin K, Tollefson AE;

XX WPI: 2001-103079/11.

XX Recombinant vector which is replication-competent in a neoplastic cell

XX and overexpresses an adenovirus death protein, useful in cancer therapy

XX when used together with replication-defective adenovirus which

XX expresses an anti-cancer gene -

XX Example 9; Fig 20: 196pp; English.

XX The invention relates to a recombinant vector (VI) which is replication-

XX competent in a neoplastic cell and which overexpresses an adenovirus

XX death protein (ADP). The vector can be used in a method for promoting

XX death of a neoplastic cell that comprises contacting the neoplastic cell

XX with at least one VI; and a composition comprising VI and a second

XX recombinant virus which is: (a) replication defective and which

XX expresses an anti-cancer gene product, where VI complements replication

XX of the second recombinant virus; or (b) replication-competent in a

XX neoplastic cell. VI, together with one or more replication-defective

XX adenovirus which expresses an anti-cancer gene product, are useful in

XX cancer therapy. Overexpression of ADP by VI results in faster lysis of

XX cells and spread of the virus throughout a cell monolayer than viruses

XX expressing wild-type levels of ADP. The present sequence represents the

XX amino acid sequence of an Ad2 ADP mutant.

XX Example 9; Fig 20: 196pp; English.

XX The invention relates to a recombinant vector (VI) which is replication-

XX competent in a neoplastic cell and which overexpresses an adenovirus

XX death protein (ADP). The vector can be used in a method for promoting

RESULT 15
AUG 61 1977

AAB61873
 ID AAB61873 standard; Protein; 40 AA.

AC AAB61873;

DT 08-MAY-2001 (first entry)

DE Ad2 ADP putative luminal domain.

KW Adenovirus death protein; ADP; neoplastic; cell death; cancer therapy;
KW Anti-cancer gene therapy; cytostatic; Ad2

XX Mastadenovirus
CS

XX
PN W0200104282-A2

XX 18-JAN-2001.
PD

12-JUL-2000:

XX 12-JUL-1999; 99US-0351778.
PR

PA (UYSL-) UNIV SAINT LOUIS.

Wold WSM, Toth K, Doronin K, Tollefson AE, PI

WPI; 2001-103079/11.

PT Recombinant vector w

PT when used together with replication-defective adenovirus which

XX Example 9: Flg 20: 196pp: English
PS

CC The invention relates to a recombinant vector (VI) which is replication
CC competent in a neoplastic cell and which overexpresses an adenovirus
CC death protein (AdD). The vector can be used in a method for promoting
CC death of a neoplastic cell that comprises contacting the neoplastic cell
CC with at least one VI; and a composition comprising VI and a second
CC recombinant virus which is: (a) replication defective and which
CC expresses an anti-cancer gene product, where VI complements replication
CC of the second recombinant virus; or (b) replication-competent in a
CC neoplastic cell. VI, together with one or more replication-defective
CC adenovirus which expresses an anti-cancer gene product, are useful in
CC cancer therapy. Overexpression of AdD by VI results in faster lysis of
CC cells and spread of the virus throughout a cell monolayer than viruses
CC expressing wild-type levels of AdD. The present sequence represents the
CC amino acid sequence of an Ad2 AdD putative luminal domain.

Sequence 40 AA;

Query Match	Score 28;	DB 22;	Length 40;
33.38;			
100.00;			

Matches	28;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
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1 MTGSTIAPTDDYRNTTATGLTSALNPQ 28

Db 1 mcgstlaptcdyrrtcatglttsalnlpq 28

Search completed: June 21, 2002, 08:23:32
Job time: 197 sec